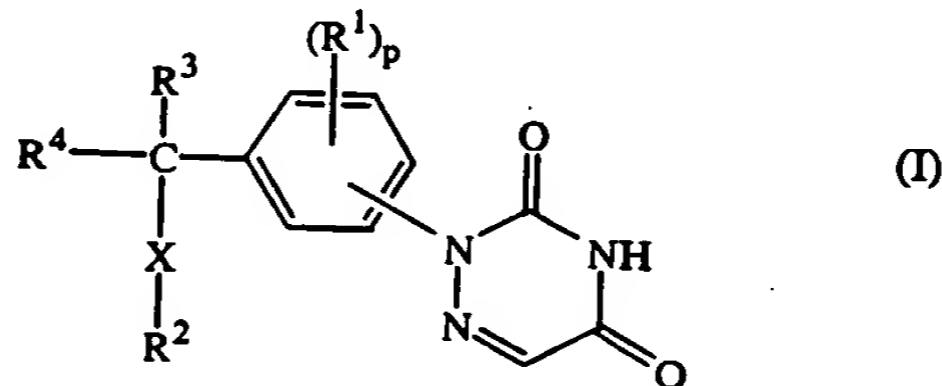


ABSTRACT

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IL-5 INHIBITING 6-AZAUACIL DERIVATIVES

The present invention is concerned with the compounds of formula



- the *N*-oxides, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein *p* is 0 to 4; *X* is O, S, NR⁵ or a direct bond; *Y* is O, S, NR⁵ or S(O)₂; R¹ independently is C₁₋₆alkyl, halo, polyhaloC₁₋₆alkyl, hydroxy, mercapto, C₁₋₆alkyloxy, C₁₋₆alkylthio, C₁₋₆alkylcarbonyloxy, aryl, cyano, nitro, Het³, R⁶, NR⁷R⁸ or substituted C₁₋₄alkyl; R² is Het¹, C₃₋₇cycloalkyl or optionally substituted C₁₋₆alkyl and if *X* is O, S or NR⁵, then R² may also represent aminocarbonyl, aminothiocarbonyl, C₁₋₄alkylcarbonyl, C₁₋₄alkylthiocarbonyl, arylcarbonyl, arylthiocarbonyl, Het¹carbonyl or Het¹thiocarbonyl; R³ and R⁴ independently are hydrogen, C₁₋₆alkyl or C₃₋₇cycloalkyl; R³ and R⁴ form a C₂₋₆alkanediyl; R⁵ is hydrogen or C₁₋₄alkyl; R⁶ is a sulfonyl or sulfinyl derivative; R⁷ and R⁸ are independently hydrogen, optionally substituted C₁₋₄alkyl, aryl, a carbonyl containing moiety, C₃₋₇cycloalkyl, -Y-C₁₋₄alkanediyl-C(=O)-O-R¹⁴, Het³, Het⁴ and R⁶; R¹¹ is hydroxy, mercapto, cyano, nitro, halo, trihalomethyl, C₁₋₄alkyloxy, formyl, trihaloC₁₋₄alkylsulfonyloxy, R⁶, NR⁷R⁸, C(=O)NR⁷R⁸, C₁₋₄alkanediyl-C(=O)-O-R¹⁴, -C(=O)-O-R¹⁴, -Y-C₁₋₄alkanediyl-C(=O)-O-R¹⁴, aryl, aryloxy, arylcarbonyl, C₃₋₇cycloalkyl, C₃₋₇cycloalkyloxy, phthalimide-2-yl, Het³ and C(=O)Het³; R¹⁴ is hydrogen, C₁₋₄alkyl, C₃₋₇cycloalkyl, aminocarbonylmethylene or mono-or di(C₁₋₄alkyl)aminocarbonylmethylene; aryl is optionally substituted phenyl; Het¹, Het², Het³ and Het⁴ are optionally substituted heterocycles; to processes for their preparation and compositions comprising them. It further relates to their use as a medicine.